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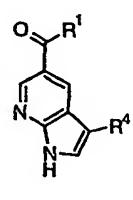
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(54) Title: 3,5-DISUBSTITUTED 1H-PZRROLO [2,3-B] PYRIDINES AS JNK INHIBITORS



**(i)** 

(57) Abstract: The present invention provides novel compounds of formula (I) and their use in the inhibition of c-Jun N-terminal kinases. The present invention further provides the use of these compounds in medicine, in particular in the prevention and/or treatment of neurodegenerative disorders related to apoptosis and/or inflammation.

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